Application No.: 10/699,635 Docket No.: IFM-005CP4CN2RCE
Group Art Unit: 1618 Examiner: F.A. Zohreh

## AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions of the claims and listing of the claims in the application;

## 1.-13. (Cancelled)

14. (Currently Amended) <u>A method for treating a subject for glaucoma, comprising administering a therapeutically effective amount of a deprenyl compound to a subject, The method of elaim 1, wherein the deprenyl compound is represented by the structure:</u>

$$R_4 = R_3 = CH = N$$
 $R_2 = CH_2 = CECH$ 
(III)

in which

R<sub>1</sub> is hydrogen, alkyl, alkenyl, alkynyl, aralkyl, alkylcarbonyl, arylcarbonyl, alkoxycarbonyl, or aryloxycarbonyl;

R2 is hydrogen or alkyl;

R<sub>3</sub> is a bond or methylene; and

R4 is aryl or aralkyl; or

R<sub>2</sub> and R<sub>4</sub>-R<sub>3</sub> are joined to form, together with the methine to which they are attached, a cyclic or polycyclic group;

and pharmaceutically acceptable salts thereof.

15. (Currently Amended) <u>A method for treating a subject for glaucoma, comprising administering a therapeutically effective amount of a deprenyl compound to a subject, The method of elaim 1, wherein the deprenyl compound is represented by the structure:</u>

$$R_4$$
— $R_3$ — $CH$ — $N$ 
 $R_2$ 
 $R_5$ — $C\equiv CH$ 
(III)

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in which

R2 is hydrogen or alkyl;

R<sub>3</sub> is a bond or methylene; and

R4 is aryl or aralkyl; or

 $R_2$  and  $R_4$ - $R_3$  are joined to form, together with the methine to which they are attached, a cyclic or polycyclic group; and

R5 is alkylene, alkenylene, alkynylene and alkoxylene;

and pharmaceutically acceptable salts thereof.

16. (Currently Amended) <u>A method for treating a subject for glaucoma, comprising administering a therapeutically effective amount of a deprenyl compound to a subject, The method of elaim 1, wherein the deprenyl compound is represented by the structure:</u>

$$CH_2$$
- $CH_2$ - $CH_N$ 
 $CH_3$ 
 $CH_2$ - $C\equiv CH$ 
 $(IV)$ 

in which

 $R_1$  is hydrogen, alkyl, alkenyl, alkynyl, aralkyl, alkylcarbonyl, arylcarbonyl, alkoxycarbonyl, or aryloxycarbonyl;

A is a substituent independently selected for each occurrence from the group consisting of halogen, hydroxyl, alkyl, alkoxyl, cyano, nitro, amino, carboxyl, -CF<sub>3</sub>, or azido;

n is 0 or an integer from 1 to 5;

and pharmaceutically acceptable salts thereof.

## 17.-20. (Cancelled)

(New) The method of claim 14, wherein said therapeutically effective amount of a deprenyl
compounds is administered with a pharmaceutically acceptable carrier.

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(New) The method of claim 21, wherein said pharmaceutically acceptable carrier has a pH
 of between about 3 and about 5

- (New) The method of claim 22, wherein said pharmaceutically acceptable carrier is an alcohol or aqueous alcohol solution.
- (New) The method of claim 21, wherein said pharmaceutically acceptable carrier is suitable for ophthalmic administration.
- 25. (New) The method of claim 14, wherein said subject is a human.
- 26. (New) The method of claim 15, wherein said therapeutically effective amount of a deprenyl compounds is administered with a pharmaceutically acceptable carrier.
- (New) The method of claim 26, wherein said pharmaceutically acceptable carrier has a pH of between about 3 and about 5.
- 28. (New) The method of claim 26, wherein said pharmaceutically acceptable carrier is an alcohol or aqueous alcohol solution.
- (New) The method of claim 28, wherein said pharmaceutically acceptable carrier is suitable for ophthalmic administration.
- 30. (New) The method of claim 15, wherein said subject is a human.
- 31. (New) The method of claim 16, wherein said therapeutically effective amount of a deprenyl compounds is administered with a pharmaceutically acceptable carrier.

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32. (New) The method of claim 31, wherein said pharmaceutically acceptable carrier has a pH of between about 3 and about 5.

- 33. (New) The method of claim 31, wherein said pharmaceutically acceptable carrier is an alcohol or aqueous alcohol solution.
- 34. (New) The method of claim 31, wherein said pharmaceutically acceptable carrier is suitable for ophthalmic administration.
- 35. (New) The method of claim 16, wherein said subject is a human.